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| APPLICATION NO. | FILING DATE | FIRST NAMED INVENTOR | ATTORNEY DOCKET NO. | CONFIRMATION NO. |
|--|-------------------------------|----------------------|---------------------|------------------|
| 10/588,028 | 04/30/2007 | Himanshu Brahmbhatt | 060348-0149 | 1320 |
| | 7590 10/06/200 LARDNER LLP | EXAMINER | | |
| SUITE 500 | | | SINGH, ANOOP KUMAR | |
| 3000 K STREET NW WASHINGTON, DC 20007 | | | ART UNIT | PAPER NUMBER |
| | | | 1632 | |
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| | | | MAIL DATE | DELIVERY MODE |
| | | | 10/06/2008 | PAPER |

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

| | Application No. | Applicant(s) | | | | |
|--|--|---|---|--|--|--|
| Office Action Commence | 10/588,028 | BRAHMBHATT ET AL. | | | | |
| Office Action Summary | Examiner | Art Unit | | | | |
| | Anoop Singh | 1632 | | | | |
| The MAILING DATE of this communication app Period for Reply | ears on the cover sheet with the c | orrespondence address | | | | |
| A SHORTENED STATUTORY PERIOD FOR REPLY WHICHEVER IS LONGER, FROM THE MAILING DA - Extensions of time may be available under the provisions of 37 CFR 1.13 after SIX (6) MONTHS from the mailing date of this communication. - If NO period for reply is specified above, the maximum statutory period w - Failure to reply within the set or extended period for reply will, by statute, Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b). | ATE OF THIS COMMUNICATION 6(a). In no event, however, may a reply be timil apply and will expire SIX (6) MONTHS from cause the application to become ABANDONEI | Lely filed the mailing date of this communication. (35 U.S.C. § 133). | | | | |
| Status | | | | | | |
| 1) Responsive to communication(s) filed on | | | | | | |
| | - action is non-final. | | | | | |
| 3) Since this application is in condition for allowan | ce except for formal matters, pro | secution as to the merits is | | | | |
| closed in accordance with the practice under <i>Ex parte Quayle</i> , 1935 C.D. 11, 453 O.G. 213. | | | | | | |
| Disposition of Claims | | | | | | |
| 4) Claim(s) <u>1-38</u> is/are pending in the application. | or form a maid matin | | | | | |
| 4a) Of the above claim(s) is/are withdraw 5) Claim(s) is/are allowed. | n from consideration. | | | | | |
| 6) Claim(s) is/are allowed. | | | | | | |
| 7) Claim(s) is/are objected to. | | | | | | |
| 8) Claim(s) <u>1-38</u> are subject to restriction and/or e | lection requirement. | | | | | |
| · · · · · · · · · · · · · · · · · · · | | | | | | |
| Application Papers | | | | | | |
| 9) The specification is objected to by the Examiner | | • | | | | |
| 10) The drawing(s) filed on is/are: a) acce | | | | | | |
| Applicant may not request that any objection to the one of Replacement drawing sheet(s) including the correction | | | | | | |
| 11) The oath or declaration is objected to by the Exa | | , , | • | | | |
| ,— | ammor. Note the attached office | 7.00.011.01.101.11.1.1.0.102. | | | | |
| Priority under 35 U.S.C. § 119 | | | | | | |
| 12) Acknowledgment is made of a claim for foreign | priority under 35 U.S.C. § 119(a) | -(d) or (f). | | | | |
| a) ☐ All b) ☐ Some * c) ☐ None of: 1. ☐ Certified copies of the priority documents | have been received | | | | | |
| 1. Certified copies of the priority documents2. Certified copies of the priority documents | | on No | | | | |
| 3. Copies of the certified copies of the prior | | | | | | |
| application from the International Bureau | | a in this National Stage | | | | |
| | * See the attached detailed Office action for a list of the certified copies not received. | | | | | |
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| Attachment(s) | ,. □ | (DTO 440) | | | | |
| 1) Notice of References Cited (PTO-892) 2) Notice of Draftsperson's Patent Drawing Review (PTO-948) | 4) ∐ Interview Summary Paper No(s)/Mail Da | | | | | |
| 3) Information Disclosure Statement(s) (PTO/SB/08) | 5) Notice of Informal P | | | | | |
| Paper No(s)/Mail Date | 6) [Other: | | | | | |

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DETAILED ACTION

Claim Interpretation: Claim 38 is directed to "use" of bacterially derived intact minicell intended for treating a disease. These claims are indefinite since the claim does not set forth any steps involved in the method/process, it is unclear what method/process applicant is intending to encompass. Additionally, because these claims fails to set forth any steps involved in the process, results in an improper definition of a process, i.e., results in a claim which is not a proper process claim. For the sake of compact prosecution, the Examiner has interpreted claim 35 to be methods of using minicell in treating disease. If Applicants do not wish for these claims to be interpreted as methods of using minicell in the treatment, Applicants are invited to amend the claims, at which point, the Examiner will determine if the amended claims fall within the elected group.

Election/Restrictions

Restriction is required under 35 U.S.C. 121 and 372.

This application contains the following inventions or groups of inventions which are not so linked as to form a single general inventive concept under PCT Rule 13.1.

In accordance with 37 CFR 1.499, applicant is required, in reply to this action, to elect a single invention to which the claims must be restricted.

Group I, claim(s) 1-7, 37, drawn to a composition comprising (i) intact minicells that contain a drug molecule and (ii) a pharmaceutically acceptable carrier therefor.

Group II, claim(s) 8-33 and 38, drawn to a drug delivery method into the cytoplasm of said mammalian cell or using bacterially derived minicell in the preparation of medicament.

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Group III, claim(s) 34-36, drawn to a method of loading minicells with a drug by creating a concentration gradient of said drug between an extracellular medium.

The inventions listed as Groups I-III do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons: In the instant case, the technical feature linking group I-III is a bacterially active minicell containing a drug molecule and a pharmaceutically carrier. Sabbadini et al (US Patent 7,183,105, dated 2/27/2007, effective filing date 5/24/2001) teach compositions comprising an intact minicell that contains a functional nucleic acid. Sabbadini et al. teach that nucleic acids of the invention can be delivered by minicells containing plasmids or expression vectors comprising sequences encoding the nucleic acids, wherein the expression constructs comprise regulatory elements operably linked to a nucleotide sequence that serves as a template for a bioactive nucleic acid (see column 7). Sabbadini et al. teach a method of delivering a biologically active compound to a cell, wherein the cell displays a ligand specifically recognized by a binding moiety, comprising contacting the cell with a minicell that displays the binding moiety, wherein the minicell comprises the biologically active compound, and wherein the contents of the minicell are delivered into the cell from a minicell bound to the cell. The binding moiety can be directed to various types of ligands. Furthermore, Sabbadini et al. teach that the ligand is a compound, composition, or moiety that is capable of being specifically bound by a binding moiety, including without limitation a receptor and an antibody (see column 34). Therefore, the instant technical feature does not contribute over prior art.

Each invention is directed to distinct goal, which comprises the use bacterium, virus or cell in order to achieve its respective and intended objective. Thus, it follows from the preceding analysis that the claimed inventions listed as Groups I-III do not relate to a single general inventive concept under PCT Rule 13.1

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because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the reasons set forth above.

1. This application contains claims directed to more than one species of the generic invention. These species are deemed to lack unity of invention because they are not so linked as to form a single general inventive concept under PCT Rule 13.1.

The species are as follows:

bispecific ligand is a polypeptide or carbohydrate or glycopeptide

Applicant is required, in reply to this action, to elect a single species to which the claims shall be restricted if no generic claim is finally held to be allowable. The reply must also identify the claims readable on the elected species, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered non-responsive unless accompanied by an election.

Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to additional species which are written in dependent form or otherwise include all the limitations of an allowed generic claim as provided by 37 CFR 1.141. If claims are added after the election, applicant must indicate which are readable upon the elected species. MPEP § 809.02(a).

The claims are deemed to correspond to the species listed above in the following manner: claim 8, and claims dependent therefrom correspond to all the species listed above.

The following claim(s) are generic: claim 8.

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The species listed above do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, the species lack the same or corresponding special technical features for the following reasons: since each disclosed species do not share a common structure feature in common with respect to their binding efficacy of minicell to mammalian cell. Thus, requirement of unity of invention is not fulfilled.

Applicant is advised that the reply to this requirement to be complete must include (i) an election of a species or invention to be examined even though the requirement may be traversed (37 CFR 1.143) and (ii) identification of the claims encompassing the elected invention.

The election of an invention or species may be made with or without traverse. To preserve a right to petition, the election must be made with traverse. If the reply does not distinctly and specifically point out supposed errors in the restriction requirement, the election shall be treated as an election without traverse.

Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a request under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

The examiner has required restriction between product and process claims. Where applicant elects claims directed to the product, and the product claims are subsequently found allowable, withdrawn process claims that depend from or otherwise require all the limitations of the allowable product claim will be considered for rejoinder. All claims directed to a nonelected process invention must require all the limitations of an allowable product claim for that process invention to be rejoined.

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In the event of rejoinder, the requirement for restriction between the product claims and the rejoined process claims will be withdrawn, and the rejoined process claims will be fully examined for patentability in accordance with 37 CFR 1.104. Thus, to be allowable, the rejoined claims must meet all criteria for patentability including the requirements of 35 U.S.C. 101, 102, 103 and 112. Until all claims to the elected product are found allowable, an otherwise proper restriction requirement between product claims and process claims may be maintained. Withdrawn process claims that are not commensurate in scope with an allowable product claim will not be rejoined. See MPEP § 821.04(b). Additionally, in order to retain the right to rejoinder in accordance with the above policy, applicant is advised that the process claims should be amended during prosecution to require the limitations of the product claims. Failure to do so may result in a loss of the right to rejoinder. Further, note that the prohibition against double patenting rejections of 35 U.S.C. 121 does not apply where the restriction requirement is withdrawn by the examiner before the patent issues. See MPEP § 804.01.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Anoop Singh whose telephone number is (571) 272-3306. The examiner can normally be reached on 9:00AM-5:30PM.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Peter Paras can be reached on (571) 272- 4517. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

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Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

Anoop Singh AU 1632

/Thaian N. Ton/ Primary Examiner, Art Unit 1632